

BIONETICS

5516 Nicholson Lane Kensington, Maryland 20795

MUTAGENIC EVALUATION OF

COMPOUND 007778189

CALCIUM SULFATE

(71-96)

SUBMITTED TO

FOOD & DRUG ADMINISTRATION
- DEPARTMENT OF HEALTH, EDUCATION AND WELFARE ROCKVILLE, MARYLAND

SUBMITTED BY

LITTON BIONETICS, INC. 5516 NICHOLSON LANE KENSINGTON, MARYLAND

APRIL 15, 1975



TABLE OF CONTENTS

		<u>Page</u>	number
EVAL	UATION SUMMARY	•••	1
I	OBJECTIVE	•••	2
II	MATERIALS	• • •	2
III	METHODS	• • •	3
IV	RESULTS SECTION.		6
	SOLUBILITY PROPERTIES OF THE TEST COMPOUND	•••	6
	TOXICITY AND DOSAGE DETERMINATIONS		7
	SUMMARY OF TEST RESULTS	• • •	8
V	INTERPRETATION OF RESULTS AND CONCLUSIONS	1	5
TABUI	LATION OF DATA	A	ppendix



EVALUATION SUMMARY

Compound 007778189, Calcium Sulfate, did not exhibit genetic activity in the series of $\underline{\text{in } \text{vitro}}$ microbial assays employed in this evaluation.

DATE:

04/15/1975

SPONSOR:

Food and Drug Administration, Contract Number 223-74-2104

SUBJECT:

Evaluation of Test Compound 007778189, Calcium Sulfate

I. OBJECTIVE

The objective of this study was to evaluate the test compound for genetic activity in microbial assays with and without the addition of mammalian metabolic activation preparations.

II. MATERIALS

A. Test Compound

Date Received:

Date Received: August, 1974

2.

Description: Fine white powder

B. <u>Indicator Microorganisms</u>

The following strains of indicator microorganisms were used in the evaluation:

Yeast Strain: Saccharomyces cerevisiae, strain D4

Bacteria Strains: Salmonella typhimurium, strains:TA-1535

TA-1537

TA-1538

C. Reaction Mixture

The following reaction mixture was employed in the activation tests:

Component Final Concentration/ml TPN (sodium salt) 6 μМ 2. Isocitric acid 49 μМ 3. Tris buffer, pH 7.4 28 μМ MgC1₂ 4. 1.7 µ M Tissue homogenate fraction 72 mg



D. <u>Tissue Homogenates and Supernatant</u>

The tissue homogenates and 9,000 x \underline{g} supernatants were prepared from tissues of the following mammalian species: Mouse-ICR random bred adult males; rat-Sprague-Dawley adult males and; primate- \underline{Macaca} mulatta adult males.

E. <u>Positive Control Compounds</u>

Table 1 lists chemicals for positive controls in the direct and activation assays.

TABLE_1 POSITIVE CONTROLS USED IN DIRECT AND ACTIVATION ASSAYS

Assay	<u>Chemical^a</u>	Solvent	Probable Mutagenic Specificity
Non-activation	Ethylmethane sulfonate	Water or saline	BPS
	2-Nitrofluorene	Dimethylsulfoxide ^C	FS
	Quinacrine mustard	Water or saline	FS
Activation	Dimethylnitrosamine	Water or saline	BPS
	2-Acetylaminofluorene	Dimethylsulfoxide ^C	FS

Concentrations given in the Results Section
BPS = base-pair substitution; FS = frameshift
Previously shown to be non-mutagenic

III. METHODS

A. <u>Toxicity</u>

The solubility, toxicity and doses for all chemicals were determined prior to screening.

Each chemical was tested for survival against the specific indicator strains over a range of doses to determine the 50% survival dose. Bacteria were tested in phosphate buffer, pH 7.4, for one hour at 37°C on a shaker. Yeasts were tested in phosphate buffer, pH 7.4, for four hours at 30°C on a shaker. The 50% survival curve and the 1/4 and 1/2 50% doses calculated.

If no toxicity was obtained for a chemical with a given strain, then a maximum dose of 5% (w/v) was used against the strain.

Unless otherwise specified, the doses calculated for the tests in buffer were applied to the activation tests. The solubility of the test chemical under treatment conditions is stated in the Results Section.



B. Plate Tests

In the nonactivation procedure, approximately 10° cells of a log-phase culture of the bacterial indicator strains were spread over the surface of a minimal plate, and a measured amount of the test chemical was placed in the center of the test plate. In activation tests, the test chemical was added to the cells, and an aliquot of the mixture was spread on the surface of the test plate. The reaction mixture (0.1 ml) plus tissue extract was then spotted on the surface of the plate. Positive and solvent controls were included. All plates were incubated at 37°C for four days and then scored. Each compound (test, positive control and solvent control) was done in duplicate. Concentrations of the positive control compounds are listed in the Results Section.

C. Suspension Tests

Non activation

Log-phase bacteria and stationary-phase yeast cultures of the indicator organisms were grown in complete broth, washed and resuspended in 0.9% saline to densities of 1 x 10^9 cells/ml and 5 x 10^7 cells/ml, respectively. This constituted the working stock for tests of a group of test chemicals and their respective controls. Tests were conducted in plastic tissue culture plates. Cells plus appropriate volume(s) of the test chemical were added to the wells to give a final volume of 1.5 ml. The solvent replaced the test chemical in the negative controls. Treatment was at 30°C for four hours for yeast tests and at 37°C for one hour for bacterial tests. All flasks were shaken during treatment. Following treatment, the plates were set on ice. Aliquots of cells were removed, diluted in sterile saline (4°C) and plated on the appropriate complete media. Undiluted samples from flasks containing the bacteria were plated on minimal selective medium in reversion experiments. from a 10^{-1} dilution of treated cells were plated on the selected media for enumeration of gene conversion with strain D4. Bacterial plates were scored after incubation for 48 hours at 37°C. The yeast plates were incubated at 30°C for 3-5 days before scoring.

2. Activation

Bacteria and yeast cells were grown and prepared as described in the non activation tests. Measured amounts of the test and control chemicals plus 0.25 ml of the stock-cell suspension were added to wells of the Linbro plate containing the appropriate tissue fraction and reaction mixture. All flasks (bacteria and yeast) were incubated at 37°C in an oxygen atmosphere with shaking. The treatment times as well as the dilutions, plating procedures and scoring of the plates were the same as described for non activation tests.



D. Preparation of Tissue Homogenates and 9,000 x g Cell Fractions

Male animals (sufficient to provide the necessary quantities tissues) were killed by cranial blow, decapitated and bled. Organs were immediately dissected from the animal using aseptic techniques and placed in ice-cold 0.25 M sucrose buffered with Tris at pH of 7.4. Upon collection of the desired quantity of organs, they were washed twice with fresh buffered sucrose and completely homogenized with a motor-driven homogenizing unit at 4° C. The whole organ homogenate obtained from this step was divided into two samples. One sample was frozen at -80° C and the other was centrifuged for 20 minutes at $9,000 \times g$ in a refrigerated centrifuge. The supernatant from the centrifuged sample was retained and frozen at -80° C. These two frozen samples were used for the activation studies.

E. Data Recording and Reporting

Following the specified incubation periods all population plates were scored by an automatic colony counter and the results from each plate of a set were recorded, in ink, on data processing forms. All minimal or other types of selective media plates were hand scored and the results recorded along with the respective population data. Other relevant experimental data were recorded on experimental definition forms. For bacteria strains the number of colonies recorded from either the population or selective plates represents that number in 1 ml of test suspension plated. The numbers recorded for the yeast strain D4 represent the number in 0.5 ml of test suspension plated. Data was then processed and printed from a computer program.

SOLUBILITY PROPERTIES OF THE TEST COMPOUND

1. NAME OR DESCRIPTION OF TEST COMPOUND:

007778189 Calcium Sulfate

2. TEST SOLVENT AND DESCRIPTION OF SOLUBILITY:

Suspension in 10% Saline Not completely soluble under treatment conditions

3. OTHER COMMENTS:

Fine white powder



TOXICITY AND DOSAGE DETERMINATIONS

COMPOUND	007778189	TEST DA	NTE: January 7,	1975

Range of concentrations of the test compound used to determine the 50% survival level

Dose Number	•	% Concentration		
1	•	10.0		
2		1.0		
3		0.1		
4		0.01		
5		0.001		

Concentrations of the test chemical required for mutagenicity tests

<u>Dose</u>	% Concentration		
	Bacteria	Yeast	
1/4 50% survival	1.25	0.625	
1/2 50% survival	2.50	1.25	
50% survival	5.00	2.50	
Plate Test	2.50		



C. <u>Summary of Test Results</u>

Plate Tests

- 1. Name or code designation of the test compound: 007778189
- 2. Test date: January 31, 1975
- 3. Concentration of the test compound: 2.5%

Test	<u>Species</u>	<u>Tissue</u>	<u>TA</u>	<u>-1535</u>	TA-1	<u>537</u>	TA.	-1538
Non-activation			1	<u>2</u>	1	2	1	<u>2</u>
Solvent Control Positive Control ^a Test Compound			6 >10 ⁴ 5	1 >10 ⁴ 3	4 85 0	3 74 0	6 34 3	2 40 0
Activation								
Negative Control Solvent Control Reaction Mixture Control			11 4	12 9	1	2 1	12 6	9 5
Positive Control Positive Control Positive Control	Mouse	Liver Lung Testes	>500 9 3	>500 > 5 4	-100 8 7	85 11 8	>200 15 10	>200 13 9
Positive Control Positive Control Positive Control	Rat	Liver Lung Testes	>100 9 4	>100 4 3	28 6 8	24 7 6	63 12 9	63 8 12
Positive Control Positive Control Positive Control	Monkey	Liver Lung Testes	>100 10 4	>100 5 5	38 6 6	25 7 6	31 12 10	28 6 10
Test Compound Test Compound Test Compound	Mouse	Liver Lung Testes	5(4(4(c) 6`	4(c) 5(c) 2	3 3 3	5 7 2	3 3 2
Test Compound Test Compound Test Compound	Rat	Liver Lung Testes	2(4(4(c) 7	3(c) 4(c) 2	3 3 3	5 7 6	3 3 2
Test Compound Test Compound Test Compound	Monkey	Liver Lung Testes	3(5(4(c) 5	5(c) 5 2	3 3 3	4 7 2	4 3 2
a TA-1535 EMS TA-1537 QM TA-1538 NF	10 μ1/p1 20 μg/p1 100 μg/p1	ate	TA	-1535 -1537 -1538	DMNA AAF AAF	10	50 μm/ 00 μg/ 00 μg/	



DATA TABLE TERMS AND ABBREVIATIONS

ABBREVIATION OR TERM	DEFINITION OR EXPLANATION
COMPOUND	Client designated compound number appears in this column.
TEST CODES	NAN = Non Activation: Solvent Control NAP = Non Activation: Positive Control NA1 = Non Activation: Test Compound Dose 1 NA2, etc. = Reflects the other dose level(s)
	A+C = Negative Chemical Control A-C = Activation: Solvent Control ACP = Activation: Positive Control ACT = Activation: Test Compound A+T = Activation: Tissue Control
	LI = Liver Tissue Activation Fraction LU = Lung Tissue Activation Fraction KI = Kidney Tissue Activation Fraction TE = Testes Tissue Activation Fraction 1,2, etc. = Dose Levels
CONCENTRATION	All test compound dose levels are expressed as a whole number followed by an exponent (negative) identified by the appropriate units.
	Example: 0025-2PCT = 0.25 percent concentration
POPU	Total number of viable cells in the plating sample raised to some exponent printed directly below the abbreviation (i.e., EP + $6 = X \cdot 10^6$).
MUT 1	Total number of mutants or convertants obtained from the sample plated raised to some exponent printed directly below the abbreviation (i.e., EP + 0 = \times 100). For strain D4, MUT 1 represents the number of ADE+ convertants.
MUT 2	Only used for strain D4 and represents the number of TRY+ convertants in the plated sample.
FREQ 1	The calculated mutation or gene conversion frequency times the negative exponent written directly below. For strain D4, FREQ 1 represents the ADE+ value.
FREQ 2	Only used for strain D4 and represents the TRY+ conversion frequency.
CONTAM	Presence of contamination on any plates.

DATA TABLE TERMS AND ABBREVIATIONS (continued)

ABBREVIATION OR TERM	DEFINITION OR EXPLANATION
AAF	2-Acetylaminofluorene
DMSO	Dimethylsulfoxide
DMN	Dimethylnitrosamine
EMS	Ethyl Methanesulfonate
QM	Quinacrine Mustard
NF	Nitrofluorene
SPECIES	Animal Strains
SPRDAW	Sprague Dawley Rats
ICRFL0	Flow ICR Random Bred Mice
RHESUS	Rhesus Monkey (<u>Macaca mulatta</u>)
MIXEDB	Dog, Mixed Breed
NEWZEA	New Zealand White Rabbit



LITTON BIOMETICS MUTAGENIC ACTIVITY SYSTEM REPORT EXR34

COMPOUND FREQUENCY SUMMARY REPORT 04/15/75

SPECIES

COMPOUND 007778189

		TA1535	TA1537	TA1538	TA1538	000004	000004
		HIS		HIS	HIS	ADF	TRY
TEST	NRG	EX-8	EX-8	EX-8		EX-5	EX-5
						· .	
NAN		1.61	12.08	5.80	4.04	2.21	3.14
ΝΔΡ		243.05	2686.46	469.44		66.33	77.39
NA1		1.45	1.35	39.18	6.53	2.18	2.70
NA2		1.20	5.00	9.81		2.01	2.97

LITTON BIONETICS MUTAGENIC ACTIVITY SYSTEM REPORT EXR34

COMPOUND FREQUENCY SUMMARY REPORT 04/15/75

SPECIFS ICRFLO COMPOUND 007778189

TEST	ORG	TA1535 HIS EX-8	TA1537 HIS EX-8	TA1538 HIS EX-8	0000D4 *DE EX-5	000004 TRY FX-5
ACT	A+C	0.93	3.59	4.92	6.30	12.91
ACT	A-C	1.04	3.21	5.77	6.38	13.81
ACT	PLI	128.68	6.71	24.04	9.48	29.12
ACT	PLU	1.00	2.19	7.37	6.67	16.81
ACT	PTE	1.63	2.90	7.02	4.73	17.40
ACT	LI1	1.99	2.64	10.56	8.10	21.03
ACT	LI2	2.40	2.72	6.12	8.16	29.53
ACT	LU1	1.84	2.45	8.03	5.02	11.62
ACT	LU2	2.06	2.64	9.74	5.61	14.42
ACT	TEI	2.17	4.53	11.71	5.01	18.66
ACT	TE2	1.95	3.39	9.88	7.71	20.89

LITTON BIONETICS MUTAGENIC ACTIVITY SYSTEM REPORT EXR34

COMPOUND FREQUENCY SUMMARY REPORT 04/15/75

SPECIES SPRDAW COMPOUND 007778189

TEST	ORG	TA1535 HIS EX-8	TA1537 HIS EX-8	TA1538 HIS EX-8	0000D4 ★ADF EX-5	0000D4 TRY EX-5
ACT	A+C	2.22	6.77	7.88	5.52	14.73
ACT	A-C	1.45	2.70	6.66	7.34	18.04
ACT	PLI	184.24	44.71	27.00	9.87	21.94
ACT	PLU	2.45	4.96	10.52	6.34	16.64
ACT	PTF	2.83	4.50	8.02	1.63	2.90
AC T	LII	0.82	10.69	4.53	7.95	16.18
ACT	L 12	0.17	6.38	9.09	4.89	17.97
ACT	LUI	0.79	2.27	1.71	5.82	12.16
ACT	LU2	2.49	0.50	9.98	6.28	15.98
ACT	TE 1	2.05	0.88	2.38	4.77	19.35
ACT	TE2	1.30	0.98	6.99	6.62	19.51

LITTON BIONETICS MUTAGENIC ACTIVITY SYSTEM REPORT EXR34

COMPOUND FREQUENCY SUMMARY REPORT 04/15/75

SPECIES RHESUS COMPOUND 007778189

		TA1535	TA1537	TA1538	000004	000004
TEST	ORG	HIS EX-8	HIS EX-8	HIS EX-8	ADE EX-5	TRY EX-5
1 13 1	UKG	4Λ ()	E V = ()	~ X = 0	(X -)	67-9
ACT	A+C	6.27	11.64	8.12	3.21	49.15
ACT	A-C	3.13	0.52	6.60	5.18	45.95
ACT	PLI	52.59	9.57	24.07	6.74	79.77
ACT	PLU	5.62	5.13	10.43	1.01	68.34
ACT	PTE	6.42	8.91	6.49	3.76	42.38
ACT	LII	2.80	3.60	3.60	5.72	73.80
ACT	~~L12	2.51	3.86	3.39	2.14	45.99
~		·				
ACT	LU1	2.06	3.29	4.40	2.32	46.53
ACT	LU2	2.82	5.32	3.29	3.21	33.17
ACT	TE 1	3.25	4.16	3.95	7.39	67.98
ΔΩΤ	TF2	3.49	10.54	5 70	2 07	54 54

٧. INTERPRETATION OF RESULTS AND CONCLUSIONS

Compound 007778189, Calcium Sulfate, was evaluated for genetic activity in a series of in vitro microbial assays with and without metabolic activation. The following results were obtained:

- A. Salmonella typhimurium
- 1. Plate tests

At a concentration of 2.5%, 007778189, was not mutagenic for any of the bacterial indicator organisms in either direct or activation plate assays.

2. Nonactivation suspension tests

The results of these tests were negative. The NAI dose of TA-1538 was high. A repeat test was conducted and the results were negative.

3. Activation suspension tests

The results of these tests were negative. The positive control results for TA-1537 and TA-1538 were lower than normal.

- В. Saccharomyces cerevisiae
- Nonactivation suspension tests 1.

The results of these tests were negative.

2. Activation suspension tests

The results of these tests were negative. The spontaneous background and the TRY locus was higher than normal which resulted in the positive control not being significantly different.

С. Conclusions

Compound 007778189, Calcium Sulfate did not exhibit genetic activity in any of the assays employed in this evaluation.

Submitted by:

David Brusick, Ph.D. Director of Genetics

APPENDIX Tabulation of Data



EXPERIMEN			22374-2104 DETECTOR TA1535	SPE	CIFS	PROJECT 02468 DATE - 04/15/75		
CUMPOHND	TFST	ORG ID	CONCENTRATION	POPU EP+6	MUT1 EP+0	FRFO1 FP-8	CONTAM	
	NAN		SALINE	1308	0021	1.61	, <u>(</u> 0.	
	МДР		EMS 0.002 %	1057	2569	243.05	0	
007778189	NAI	•	0025-1 PCT.	0830	0012	1.45	2	
007778189	NA2	,	0125-2 PCT.	1083	0013	1.20	?	

EXPERIMENT 502302			22374-2104 DETECTOR TA1537	SPECIES		PROJECT 02468 DATE - 04/15/75	
COMPOUND	TEST	ORG ID	CONCENTRATION	ROPH EP+6	MUT1 EP+0	FREO1	CONTAM
	ΝΔΝ		SALINE	0240	0029	12.08	0
	NAP		OM 1.0 UG/ML	0096	2579	2686.46	n
007778189	NA1		0025-1 PCT.	0668	0009	1.35	2
007778189	NA?		0125-2 PCT.	0360	0018	5.00	2

<u> </u>			22374-2104 DETECTOR TA1538	S SPE	CIFS	PROJECT 02468 DATE - 04/15/75		
COMPOUND	TEST	ORG ID	CONCENTRATION	POPU EP+6	MUT1 EP+0	FREQ1 EP-8	CONTAM	
	NAN		DMSO	0414	0024	5.80	0	
	NAP		NF 125 UG-ML	0288	1352	469.44	0	
007778189	NAI	· e	0025-1 PCT.	0342	0134	39.18	2	
007778189	NA2		0125-2 PCT.	0367	0036	9.81	0	

EXPERIMENT 505005		22374-2104 DETECTOR TA1538	SPECIES		PROJECT 02468 DATE - 04/15/75		
COMPOUND :	TECT	ORG	20112	PUPII	MUTI	FREQ1	
Characterial	TEST	ID	CONCENTRATION	EP+6	EP+0	FP-8	CONTAM
•	NAN		DMSO	0570	0023	4.04	0
007778189	NA1		0025-1 PCT.	0567	0037	6.53	0

CONTRACT EXPERIMENT 504801		22374-2104	PROJECT 02468						
		01	DETECTOR 000004	SPECIES			D	DATE - 04/15/75	
		ORG		POPII	MUT1	MUT2	FREQI	FRFQ2	
COMPOUND	TEST	ID	CONCENTRATION	FP+4	EP+1	EP+1	EP-5	FP-5	CUNTAM
	NAN		SALINE	1084	0024	0034	2.21	3.14	0
	NAP		FMS 1.0 %	1004	0666	0777	66.33	77.39	0
007778189	NAI		0125-2 PCT.	0962	0021	0026	2.18	2.70	2
007778189	NA2		0625-3 PCT.	1144	0023	0034	2.01	2.97	, 6

REPORT EXR33 LITTON BIONETICS MUTAGENIC ACTIVITY SYSTEM COMPOUND SUMMARY BACKUP DETAIL

CONTRACT 22374-2104		22374-2104	PROJECT 02468						
EXPERIMEN.	T 4346	501	DETECTOR TA1539	5 SPE	CIES	ICRFLO	DATE - 0	4/15/75	
		NRG		POPU	MUTI	FRE01			
CUMPOUND	TEST	ID	CONCENTRATION	EP+6	EP+0	EP-8		CONTAM	
* .	A+C		DMN 50 UM/ML	1286	0012	0.93		. 0	
	A-C		SALINE	1248	0013	1.04	• ·	?	
•	ACP	LI	DMN 50 UM/ML	1210	1557	128.68		0	
	ACP	, L. U	DMN 50 UM/ML	1095	.0011	1.00		2	
	ACP	TE	DMN 50 UM/ML	1163	0019	1.63	•	2	
007778189	ACT	LII	0025-1 PCT.	1204	0024	1.99		2	
007778189	ACT	ris	0125-2 PCT.	1376	0033	2.40		3	
007778189	ACT	LU1	0025-1 PCT.	1032	0019	1.84		2	
007778189	ACT	LU2	0125-2 PCT.	0971	0020	2.06	•	. 2	
007778189	ACT	TE1	0025-1 PCT.	1061	0023	2.17		2	
007778189	ACT	TE2	0125-2 PCT.	1491	0029	1.95		. 2	

CONTRACT			22374-210	14	PROJECT 02468						
EXPERIMENT	4347	01	DETECTOR	TA1537	SPE	CIFS I	CRFLO	Đ	ATE - 04	/15/75	
		ORG			POPII	MUT1		FREQ1			
COMPOUND	TEST	ID	CONCENTRA	MOITA	EP+6	EP+0		EP-8		CONTAM	
	A+C		AAF 800 U	IG/ML	1783	0064		3.59		0	
•	V -C		DMSO		1590	0051	•	3.21		n	
•	ACP	LI	AAF 800 U	IG/ML	1983	0133		6.71		3	
	ACP	LU	AAF 800 I	IG/ML	1460	0032		2.19		2	
	ACP	ΤE	AAF 800 U	IG/ML	1619	0047		2.90	•	2	
007778189	ACT	LI1	0025-1 PC	ст.	1592	0042		2.64		· 2	
007778189	ACT	LI2	0125-2 PC	Τ.	1473	0040		2.72		2	
007778189	ACT	LU1	0025-1 PC	т.	1470	0036		2.45		2	
007778189	ACT	LU2	0125-2 PC	Τ.	1404	0037		2.64		2	
007778189	ACT	TE1	0025-1 PC	т.	1765	0080		4.53		2	
007778189	ACT.	TF2	0125-2 PC	T	1623	0055		3.39		2	

•		22374-2104	PROJECT 02468						
EXPERIMENT	T 4350	001	DETECTOR TA1538	SPF	SPECIES ICRELO DATE - 04/15/				
		ORG		POPII	MUTI	FREQI			
COMPOUND	TEST	ID	CONCENTRATION	EP+6	EP+0	FP-8	CONTAM		
	A+C		AAF 800 UG/ML	1363	0067	4.92	n		
•	A-C		DMSO	1820	0105	5.77	0		
	ACP	LI	AAF 800 UG/ML	1252	0301	24.04	3		
	ACP	LU	AAF 800 UG/ML	1262	0093	7.37	. 2		
	ACP	TE	AAF 800 UG/ML	1140	0080	7.02	.		
007778189	ACT	t. I 1	0025-1 PCT.	1023	0108	10.56	2		
007778189	ACT	LIZ	0125-2 PCT.	1144	0070	6.12	2		
007778189	ACT	LUI	0025-1 PCT.	1195	0096	8.03	2		
007778189	ACT	LU2	0125-2 PCT.	0955	0093	9.74	2		
007778189	ACT	TE1	0025-1 PCT.	0743	0087	11.71	· · · 2		
007778189	ACT	TE2	0125-2 PCT.	1042	0103	9.88	2		

	22374-21	04	PROJECT 02468							
EXPERIMENT	F 5007	01	DETECTOR	000004	SPE	CIES	ICRFLO	'nD	ATE - 04/	15/75
		ORG		1	POPU	MUT1	MUT2	FREQ1	FRE02	
CUMPOUND	TEST	ID	CONCENTR	ATION	EP+4	EP+1	EP+1	FP-5	EP-5	CUNTAM
	A+C		DMN 90 U	M/ML	0968	0061	0125	6.30	12.91	0
•	A-C		SALINE	•	1050	0067	0145	6.38	13.81	0
	ACP	LI	DMN 90 U	M/ML	0728	0069	0212	9.48	29.12	2
	ACP	LU	DMN 90 U	M/ML	0809	0054	0136	6.67	16.81	2
	ACP	TE	DMN 90 U	M/MI_	0931	0044	0162	4.73	17440	6
007778189	ACT	LII	0125-2 P	CT.	0951	0077	0200	8.10	21.03	6
007778189	AC T	LIZ	0625-3 PC	CT.	0674	0055	0199	8.16	29.53	6
007778189	ACT	LU1	0125-2 P	CT.	0697	0035	0081	5.02	11.62	6
007778189	ACT	LU2	0625-3 PC	CT.	0874	0049	0126	5.61	14.42	6
007778189	ACT	TE1	0125-2 P	CT.	0659	0033	0123	5.01	18.66	6
007778189	AÇ T	TE2	0625-3 PC	CT.	0895	0069	0187	7.71	20.89	3

REPORT EXR33 LITTON BIONETICS MUTAGENIC ACTIVITY SYSTEM COMPOUND SUMMARY BACKUP DETAIL

			22374-2104		PROJECT 02468					
EXPERIMENT	r 5002	201	DETECTOR TA153	SPF	CIES		- 04/15/75			
		ORG		PNPU	MUT1	FRE01				
COMPOUND	TEST	ID	CONCENTRATION	EP+6	FP+0	EP-8	CONTAM			
	A+C		DMN 50 UM/ML	0450	0010	2.22	o			
	A-C		SALINE	0826	0012	1.45	2			
	ACP	LI	DMN 50 UM/ML	0628	1157	184.24	n			
	ACP	L U	DMN 50 UM/ML	0653	0016	2.45	0			
· • • • • • • • • • • • • • • • • • • •	ACP	TE	DMN 50 UM/ML	0566	0016	2.83	, 2			
007778189	ACT	LII	0025-1 PCT.	0734	0006	0.82	2			
007778189	AC T	LI2	0125-2 PCT+	0590.	0001	0.17	2			
007778189	ACT	LU1	0025-1 PCT.	0632	0005	0.79	2			
007778189	ACT	LII2	0125-2 PCT.	0442	0011	2.49	2			
007778189	ACT	TF1	0025-1 PCT.	0584	0012	2.05	3			
007778189	AC T	TE2,	0125-2 PCT.	0540	0007	1.30	2			

	TRACT	22374-2104		PROJECT 02468					
EXPERIMENT	T 5021	01	DETECTOR TA	1537	SPE	CIFS	SPRDAW	DATE -	04/15/75
		ORG		į	POPII	MUT1	FREO	1	
CUMBUNND	TEST	ID	CONCENTRATI	UN	EP+6	FP+0	FP-8		CONTAM
	A+C		AAF 800 UG/	MI_ (0251	0017	6.7	7 .	0
	A-C		DMSO		0185	0005	2.7	n _.	n
	ACP	L I	AAF 800 UG/	ML (0170	0076	44.7	1 .	2
	ACP	LU	AAF 800 UG/	ML	0141	0007	4.9	6	n
	ACP	TE	AAF 800 UG/	ML (0111	0005	4.5	•	0
007778189	ACT	LII	0025-1 PCT.		0533	0057	10.6	9	2
007778189	ACT	LI2	0125-2 PCT.		0893	0057	6.38	3	2
007778189	ACT	LU1	0025-1 PCT.		0088	0002	2.2	7	2
007778189	ACT	LU2	0125-2 PCT.	(0201	0001	0.50		2
007778189	ACT	TE1	0025-1 PCT.	ı	0452	0004	0.8	8	2
007778189	AC T	TE2	0125-2 PCT.	(307	0003	0.98	3	2

REPORT EXR33 LITTON BIONETICS MUTAGENIC ACTIVITY SYSTEM COMPOUND SUMMARY BACKUP DETAIL

			22374-2104	PROJECT 02468					
EXPERIMEN.	T 5003	301	DETECTOR TA1538	SPE	CIES SPRDAW	DAT	E - 04/15/75		
		ORG		POPU	MUTI	FRE01			
COMPOUND	TEST	1D	CONCENTRATION	EP+6	EP+0	EP-8	CONTAM		
	A+C		AAF 800 UG/ML	0964	0076 .	7.88	0		
	. A-C		DMSU	1276	0085	6.66	. 1		
	ACP	LI	AAF 800 UG/ML	1052	0284	27.00	. 0		
	ACP	LU	MAE 800 HG/ML	1017	0107	10.52	, o		
	ACP	TE	AAF 800 UG/ML	1,347	0108	8.02	. 2		
007778189	ACT	LII	0025-1 PCT.	0640	0029	4.53	2		
007778189	ACT	L I 2	0125-2 PCT.	1331	0121	9.09	2		
007778189	ACT	LU1	0025-1 PCT.	1990	0034	1.71	2		
007778189	ACT	LU2	0125-2 PCT.	0932	0093	9.98	2		
007778189	ACT	TF1	0025-1 PCT.	0797	0019	2.38	2		
007778189	ACT	TF2	0125-2 PCT.	1016	0071	6.99	2		

REPORT FXR33 LITTON BIONETICS MUTAGENIC ACTIVITY SYSTEM COMPOUND SUMMARY BACKUP DETAIL

		COM	TRACT	22374-2104 PROJECT 02468							
EXPERIMEN		T 500801		DETECTOR (0000D4	• ŞPE	SPECIFS SPRDAW			DATE - 04/15/75		
	•		ORG	•	POPU	MUTI	MUT2	FREQ1	FREQ2		
	COMPOUND	TEST	TD.	CONCENTRATION	EP+4	EP+1	EP+1	FP-5	EP-5	CONTAM	
		A+C		DMN 90 UM/ML	0706	0039	0104	5.52	14.73	0 -	
		A-C		SALINE	0654	0048	0118	7.34	18.04	0	
	`	ACP	LI	DMN 90. UM/ML	0679	0067	0149	9.87	21.94	0	
		ACP	LU	DMN 90 UM/ML	0631	0040	0105	6.34	16.64	o O	
		ACP	TE	DMN 90 UM/ML	0861	0014	0025	1.63	2 490	0	
	007778189	ACT	LII	0125-2 PCT.	0692	0055	0112	7.95	16.18	6	
	007778189	ACT	LIZ	0625-3 PCT.	0818	0040	0147	4.89	17.97	0 .	
	007778189	ACT	LU1	0125-2 PCT.	0773	0045	0094	5.82	12.16	6	
	007778189	AÇ T	LU2	0625-3 PCT.	0732	0046	0117	6.28	15.98	6	
	007778189	ACT	TF1	0125-2 PCT.	0796	0038	0154	4.77	19.35	6	
	007778189	AC T	TE2	0625-3 PCT.	0574	0038	0112	6.62	19.51	6	

			22374-2104	PROJECT 02468					
EXPERIMENT	T 5009	901	DETECTOR TA153	5 SPF	FCIES RHESUS	DATE	DATE - 04/15		
6 0 1 1 0 1 1 1 1		ORG		POPU	MUTI	FREO1			
CUMPOUND	TEST	ID	CONCENTRATION	EP+6	EP+O	FP-8		CONTAM	
	A+C		DMN 50 UM/ML	0734	0046	6.27		0	
	A-C .		SALINE	1119	0035	3.13		2	
	ACP	LI	DMN 50 UM/ML	0945	0497	52.59		3 .	
	ACP	LU	DMN 50 UM/ML	0908	0051	5.62		0	
	ACP	TE	DMN 50 UM/ML	0748	0048	6.42	•	2	
007778189	ACT	LI1	0025-1 PCT.	0999	0028	2.80	·	2	
007778189	ACT	L I 2	0125-2 PCT.	0917	0023	2.51		, 2	
007778189	ACT	1.01	0025-1 PCT.	1357	0028	2.06		2	
007778189	ACT	1.112	0125-2 PCT.	1099	0031	2.82		2	
007778189	ACT	TE1	0025-1 PCT.	0923	0030	3.25		2	
007778189	ACT	TF2	0125-2 PCT.	1175	0041	3 40		`a	

REPORT EXR33 LITTON BIONETICS MUTAGENIC ACTIVITY SYSTEM COMPOUND SUMMARY BACKUP DETAIL

			22374-2104	PROJECT 02468						
EXPERIMENT 502201			DETECTOR TA1537	SPF	CIFS	RHESUS	DATE - 04/15/75			
		ORG		PNPU	MUT1	FREQ1				
COMPOUND	TEST	ID	CONCENTRATION	EP+6	EP+0	FP-8		CONTAM		
	A+C		AAÉ 800 UG/ML	0146	0017	. 11.64	•	0		
,	A-C		DMSO	0192	0001	0.52)	'n		
	ACP	ΓI	AAF 800 HG/ML	0188	0018	9.57	•	n		
*	ACP	LU	AAF 800 HG/ML	0156	0008	5.13		2		
•	ACP	TE	AAF 800 UG/ML	0101	0009	8.91		• 0		
007778189	ACT	LII	0025-1 PCT.	0694	0025	3.60)	2		
007778189	ACT	1.12	0125-2 PCT.	0440	0017	3.86		2		
007778189	ACT	1.01	0025-1 PCT.	0577	0019	3.29		2		
007778189	ACT	1.112	0125-2 PCT.	0677	0036	5.32		2		
007778189	ACT	TE1	0025-1 PCT.	0529	0022	4.16		3		
007778189	ACT	TE2	0125-2 PCT.	0408	0043	10.54	`	2		

CONTRACT 22374-2104					PROJECT 02468					
EXPERIMEN:			DETECTOR TA1538	SPE	CIES RHESUS		DATE -	04/15/	75	
		ORG		POPU	MUTI	FRE01				
CUMPOUND	TEST	ID	CONCENTRATION	EP+6	FP+O	F.P-8		С	ONTAM	
	A+C		AAF 800 UG/ML	0936	0076	8.12			0	
٠	A-C		DMSO	1076	0071	6.60)		2 -	
•	ACP	LI	AAF 800 UG/ML	0810	0195	24.07			3	
	ACP	LU	AAF 800 UG/ML	1064	0111	10.43	3		0	
	ACP	TE	AAF 800 UG/ML	1263	0082	6.49)	•	2	
007778189	ACT	LII	0025-1 PCT.	0749	0027	3.60)		2	
007778189	ACT	LI2	0125-2 PCT.	1091	0037	3.39			2	
007778189	ACT	LUI	0025-1 PCT.	0796	0035	4.40) .	,	3	
007778189	ACT	LU2	0125-2 PCT.	1215	0040	3.29)		? `	
007778189	ACT	TE1	0025-1 PCT.	0759	0030	3.9	5		2	
007778189	ACT	TF2	0125-2 PCT.	1054	0061	5.79)		2	

REPORT EXR33 LITTON BIONETICS MUTAGENIC ACTIVITY SYSTEM COMPOUND SUMMARY BACKUP DETAIL

			22374-2104		PROJECT 02468						
EXPERIMENT 502901			DETECTOR 00000	4 SPI	SPECIES RHESUS			DATE - 04/15/75			
		ORG		BOPU	MUTI	MUT2	FRE01	FRE02			
COMPOUND	TFST	ID	CONCENTRATION	EP+4	EP+1	FP+1	EP-5	EP-5	CONTAM		
	A+C		DWN 90 UM/MI	0529	0017	0260	3.21	49.15	4		
·	A-C		SALINE	0618	0032	0284	5.18	45.95	0		
•	ACP	LΙ	DMN 90 UM/ML	0341	0023	0272	6.74	79.77	0		
	ACP	LU	DMN 90 UM/ML	0398	0004	0272	1.01	68.34	2		
	ACP	TE	DMN 90 UM/ML	0505	0019	0214	3.76	42.38	n		
007778189	ACT	LII	0125-2 PCT.	0332	0019	0245	5.72	73.80	6		
007778189	ACT	112	0625-3 PCT.	0561	0012	0258	2.14	45.99	2		
007778189	ACT	LUI	0125-2 PCT.	0475	0,011	0221	2.32	46.53	6		
007778189	AC T	LU2	0625-3 PCT.	0624	0020	0207	3.21	33.17	4		
007778189	ACT	TE1	0125-2 PCT.	0406	0030	0276	7.39	67.98	6		
007778189	ACT	TE2	0625-3 PCT.	0488	0015	0276	3.07	56.56	6		